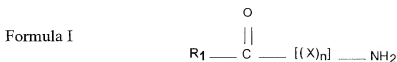


We claim:

1. An antimicrobial composition comprising a plurality of hexapeptides wherein for each hexapeptide, the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;  
the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;  
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and  
wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.
2. The antimicrobial composition of claim 1 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.
3. The antimicrobial composition of claim 1 wherein said peptides are incorporated into a polymer.
4. The antimicrobial composition of claim 3 wherein said polymer is selected from the group consisting of a polysaccharide, a glycol polymer, a polyester, a polyurethane, a polyacrylate, a polyacrylonitrile, a polyamide, a polyolefin, a polystyrene, a vinyl polymer, a polypropylene, silk, a biopolymer, and mixtures thereof.
5. An antimicrobial composition comprising a plurality of peptides, wherein said peptides

each are represented by Formula I:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>3</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>6</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkylthio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

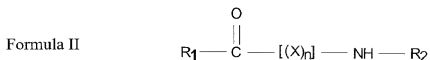
6. The antimicrobial composition of claim 5 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

7. The antimicrobial composition of claim 5 wherein said peptides are incorporated into a polymer.

8. The antimicrobial composition of claim 7 wherein said polymer is selected from the group consisting of a polysaccharide, a glycol polymer, a polyester, a polyurethane, a polyacrylate, a polyacrylonitrile, a polyamide, a polyolefin, a polystyrene, a vinyl polymer, a polypropylene, silk, a biopolymer, and mixtures thereof.

9. An antimicrobial composition comprising a plurality of peptides, wherein said peptides

each are represented by Formula II:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfanylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>2</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfanylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or

quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>8</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkythio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid.

10. The antimicrobial composition of claim 9 wherein the amino acids in the first and second positions of said peptide, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

11. The antimicrobial composition of claim 9 wherein said peptides are incorporated into a

polymer.

12. The antimicrobial composition of claim 11 wherein said polymer is selected from the group consisting of a polysaccharide, a glycol polymer, a polyester, a polyurethane, a polyacrylate, a polyacrylonitrile, a polyamide, a polyolefin, a polystyrene, a vinyl polymer, a polypropylene, silk, a biopolymer, and mixtures thereof.

13. An antimicrobial composition comprising a plurality of hexapeptides and at least one carrier, wherein for each hexapeptide:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;

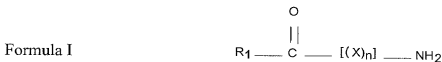
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

14. The antimicrobial composition of claim 13 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

15. The antimicrobial composition of claim 13 wherein said carrier is selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

16. An antimicrobial composition comprising a plurality of hexapeptides and at least one carrier, wherein said each hexapeptide is represented by Formula I:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>8</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkylthio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;  
and

R<sub>7</sub> is independently halogen;

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano;  
wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

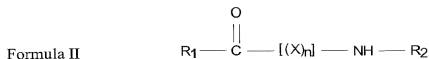
wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

17. The antimicrobial composition of claim 16 wherein the amino acids in the first and second positions of said peptides, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

18. The antimicrobial composition of claim 16 wherein said carrier is selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

19. An antimicrobial composition comprising a plurality of hexapeptides and at least one carrier, wherein said each hexapeptide is represented by Formula II:





wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfanylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>2</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfanylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>5</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>5</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>8</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>5</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkythio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano.

20. The antimicrobial composition of claim 19 wherein the amino acids in the first and second positions, based on numbered amino acids from N-terminus to C-terminus, are selected from the group consisting of Arg-Tyr, Arg-Cys, Ser-Thr, Met-Trp, Lys-Trp, Thr-Trp, Trp-Arg, Trp-His, and Trp-Tyr.

21. The antimicrobial composition of claim 19 wherein said carrier is selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

22. A method for preventing, inhibiting, or terminating the growth of at least one microbe comprising administering an antimicrobial amount of a plurality of hexapeptides and at least one carrier, wherein for each hexapeptide:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-

terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;

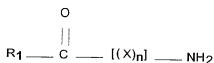
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

23. The method of claim 22 wherein said microbe comprises *Burkholderia cepacia*.

24. A method for preventing, inhibiting, or terminating the growth of at least one microbe comprising administering an antimicrobial amount of a plurality of peptides and at least one carrier, wherein said peptides are each represented by Formula I:

Formula I



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or

quinolinyl each optionally substituted with  $R_5$ ,  $R_6$  or  $R_7$ ;

$R_2$  is  $C_1$ - $C_{20}$  alkyl;  $C_3$ - $C_6$  cycloalkyl;  $C_4$ - $C_{20}$  alkenyl;  $C_4$ - $C_{20}$  alkynyl;  $C_1$ - $C_{20}$  haloalkyl;  $C_3$ - $C_{20}$  haloalkenyl;  $C_3$ - $C_{20}$  haloalkynyl;  $C_2$ - $C_{20}$  alkoxyalkyl;  $C_2$ - $C_{20}$  alkylthioalkyl;  $C_2$ - $C_{20}$  alkylsulfanylalkyl;  $C_2$ - $C_{20}$  alkylsulfonylalkyl;  $C_5$ - $C_{20}$  cycloalkylalkyl;  $C_4$ - $C_{20}$  alkenyloxyalkyl;  $C_4$ - $C_{20}$  alkynyloxyalkyl;  $C_4$ - $C_{20}$  (cycloalkyl) oxyalkyl;  $C_4$ - $C_{20}$  alkenylthioalkyl;  $C_4$ - $C_{20}$  alkynylthioalkyl;  $C_6$ - $C_{20}$  (cycloalkyl) thioalkyl;  $C_2$ - $C_{20}$  haloalkoxyalkyl;  $C_4$ - $C_{20}$  haloalkenyloxyalkyl;  $C_4$ - $C_{20}$  haloalkynyloxyalkyl;  $C_4$ - $C_{20}$  alkoxyalkenyl;  $C_4$ - $C_{20}$  alkoxyalkynyl;  $C_4$ - $C_{20}$  alkylthioalkenyl;  $C_4$ - $C_{20}$  alkylthioalkynyl;  $C_4$ - $C_{20}$  trialkylsilylalkyl;  $C_1$ - $C_{20}$  alkyl substituted with  $NR_3R_4$ , nitro, cyano, or phenyl optionally substituted with  $R_5$ ,  $R_6$ , and  $R_7$ ;  $C_1$ - $C_{20}$  alkoxy;  $C_1$ - $C_{20}$  haloalkoxy;  $C_1$ - $C_{20}$  alkylthio;  $C_1$ - $C_{20}$  haloalkylthio;  $NR_3R_4$ ; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with  $R_5$ ,  $R_6$  or  $R_7$ ;

$R_3$  is independently hydrogen;  $C_1$ - $C_4$  alkyl; or phenyl optionally substituted with at least one  $R_8$ ;

$R_4$  is independently hydrogen;  $C_1$ - $C_8$  alkyl; or phenyl optionally substituted with at least one  $R_8$ ;

$R_5$  is independently  $C_1$ - $C_6$  alkyl;  $C_1$ - $C_6$  alkoxy;  $C_1$ - $C_6$  haloalkyl; halogen;  $C_2$ - $C_8$  alkynyl;  $C_1$ - $C_6$  thioalkyl; phenyl or phenoxy each optionally substituted with at least one  $R_8$ ; cyano; nitro;  $C_1$ - $C_6$  haloalkoxy;  $C_1$ - $C_6$  haloalkylthio;  $C_2$ - $C_6$  alkenyl;  $C_2$ - $C_6$  haloalkenyl; acetyl;  $CO_2CH_3$ ; or  $N(C_1-C_2 \text{ alkyl})_2$ ;

$R_6$  is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

$R_7$  is independently halogen; and

$R_8$  is independently halogen;  $C_1$ - $C_4$  alkyl;  $C_1$ - $C_4$  alkoxy;  $C_1$ - $C_4$  haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-

terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

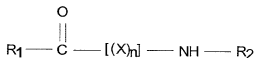
the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

25. The method of claim 24 wherein said microbe comprises *Burkholderia cepacia*.

26. A method for preventing, inhibiting, or terminating the growth of at least one microbe comprising administering an antimicrobial amount of a plurality of peptides and at least one carrier, wherein said peptides are each represented by Formula II:

Formula II



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>3</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothieryl, or

quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>2</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>8</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkylthio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano.

27. The method of claim 26 wherein said microbe comprises *Burkholderia cepacia*.

28. A composition for coating a substrate comprising an antimicrobial amount of a plurality of hexapeptides and at least one carrier, wherein for each hexapeptide:

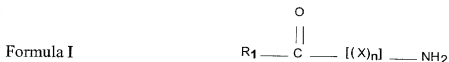
the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan;

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid; and

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

29. A composition for coating a substrate comprising an antimicrobial amount of a plurality of peptides and at least one carrier, wherein each of said peptides are represented by Formula I:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>3</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-

C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>2</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfanylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>8</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkylthio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to



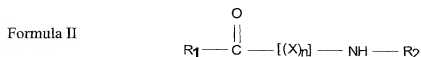
C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine-arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

30. A composition for coating a substrate comprising an antimicrobial amount of a plurality of peptides and at least one carrier, wherein each of said peptides are represented by Formula II:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 6;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or

phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>2</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

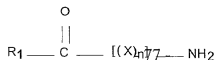
R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>8</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkylthio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano.

31. An antimicrobial composition comprising a plurality of peptides, wherein said peptides each are represented by Formula I:



## Formula I

wherein:

X represents any amino acid except glutamate or aspartate;

n = 1-10;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>6</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkylthio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano;

wherein:

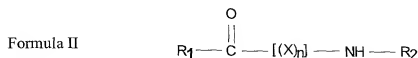
the amino acid in the first position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid;

wherein the first two amino acids of said hexapeptides are other than arginine arginine, tryptophan-tryptophan, tryptophan-cysteine, tryptophan-lysine, arginine-tryptophan, or threonine-arginine.

32. An antimicrobial composition comprising a plurality of peptides, wherein said peptides each are represented by Formula II:



wherein:

X represents any amino acid except glutamate or aspartate;

n = 1-10;

R<sub>1</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfinylalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonylalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-

C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>2</sub> is C<sub>1</sub>-C<sub>20</sub> alkyl; C<sub>3</sub>-C<sub>6</sub> cycloalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyl; C<sub>4</sub>-C<sub>20</sub> alkynyl; C<sub>1</sub>-C<sub>20</sub> haloalkyl; C<sub>3</sub>-C<sub>20</sub> haloalkenyl; C<sub>3</sub>-C<sub>20</sub> haloalkynyl; C<sub>2</sub>-C<sub>20</sub> alkoxyalkyl; C<sub>2</sub>-C<sub>20</sub> alkylthioalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonalkyl; C<sub>2</sub>-C<sub>20</sub> alkylsulfonalkyl; C<sub>5</sub>-C<sub>20</sub> cycloalkylalkyl; C<sub>4</sub>-C<sub>20</sub> alkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> (cycloalkyl) oxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkenylthioalkyl; C<sub>4</sub>-C<sub>20</sub> alkynylthioalkyl; C<sub>6</sub>-C<sub>20</sub> (cycloalkyl) thioalkyl; C<sub>2</sub>-C<sub>20</sub> haloalkoxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkenyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> haloalkynyloxyalkyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkenyl; C<sub>4</sub>-C<sub>20</sub> alkoxyalkynyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkenyl; C<sub>4</sub>-C<sub>20</sub> alkylthioalkynyl; C<sub>4</sub>-C<sub>20</sub> trialkylsilylalkyl; C<sub>1</sub>-C<sub>20</sub> alkyl substituted with NR<sub>3</sub>R<sub>4</sub>, nitro, cyano, or phenyl optionally substituted with R<sub>5</sub>, R<sub>6</sub>, and R<sub>7</sub>; C<sub>1</sub>-C<sub>20</sub> alkoxy; C<sub>1</sub>-C<sub>20</sub> haloalkoxy; C<sub>1</sub>-C<sub>20</sub> alkylthio; C<sub>1</sub>-C<sub>20</sub> haloalkylthio; NR<sub>3</sub>R<sub>4</sub>; or phenyl, benzyl, pyridyl, furanyl, thienyl, naphthyl, pyrimidinyl, benzofuranyl, benzothienyl, or quinolinyl each optionally substituted with R<sub>5</sub>, R<sub>6</sub> or R<sub>7</sub>;

R<sub>3</sub> is independently hydrogen; C<sub>1</sub>-C<sub>4</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>4</sub> is independently hydrogen; C<sub>1</sub>-C<sub>8</sub> alkyl; or phenyl optionally substituted with at least one R<sub>8</sub>;

R<sub>5</sub> is independently C<sub>1</sub>-C<sub>6</sub> alkyl; C<sub>1</sub>-C<sub>6</sub> alkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkyl; halogen; C<sub>2</sub>-C<sub>8</sub> alkynyl; C<sub>1</sub>-C<sub>6</sub> thioalkyl; phenyl or phenoxy each optionally substituted with at least one R<sub>8</sub>; cyano; nitro; C<sub>1</sub>-C<sub>6</sub> haloalkoxy; C<sub>1</sub>-C<sub>6</sub> haloalkylthio; C<sub>2</sub>-C<sub>6</sub> alkenyl; C<sub>2</sub>-C<sub>6</sub> haloalkenyl; acetyl; CO<sub>2</sub>CH<sub>3</sub>; or N(C<sub>1</sub>-C<sub>2</sub> alkyl)<sub>2</sub>;

R<sub>6</sub> is independently methyl; ethyl; methoxy; methylthio; halogen; or trifluoromethyl;

R<sub>7</sub> is independently halogen; and

R<sub>8</sub> is independently halogen; C<sub>1</sub>-C<sub>4</sub> alkyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> haloalkyl; nitro; or cyano;

wherein:

the amino acid in the first position, based on numbered amino acids from N-terminus to

C-terminus, is selected from the group consisting of arginine, lysine, methionine, serine, threonine and tryptophan;

the amino acid in the second position, based on numbered amino acids from N-terminus to C-terminus, is selected from the group consisting of arginine, histidine, cysteine, threonine, tyrosine, and tryptophan; and

the amino acids in positions three through six, based on numbered amino acids from N-terminus to C-terminus, are any amino acid.

33. The antimicrobial composition of claim 31 further comprising a carrier selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.

34. The antimicrobial composition of claim 32 further comprising a carrier selected from the group consisting of a pharmaceutically acceptable carrier, an industrially acceptable carrier, a household product, and a personal care composition.